AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1.-28. (Canceled)

29. (Currently Amended) A method for identifying a compound that modulates the activity of a human c-Maf protein, comprising

contacting the indicator composition with a test compound; and

providing an indicator composition cell that comprises a human c-Maf protein of SEQ ID NO.:2, and a reporter gene responsive to the human c-Maf protein;

determining the effect of the test compound on the activity of the human c-Maf protein in the indicator composition cell wherein the step of determining comprises evaluating the expression of the reporter gene in the presence and absence of the test compound, to thereby identify a compound that modulates the activity of a human c-Maf protein.

30.-31. (Cancel)

32. (Currently Amended) The A method for identifying a compound that modulates an immune response, comprising of claim 29, further comprising providing an indicator cell that comprises a human c-Maf protein of SEQ ID

NO.:2, and a Th2-associated cytokine gene responsive to the human c-Maf protein;

contacting the indicator composition with a test compound; and

determining the effect of the test compound on an immune response, wherein the step of determining comprises evaluating the effect of the compound on expression of the Th2-associated cytokine gene in the presence and the absence of the test compound, to thereby identify a compound that modulates an immune response.

33. (Cancel)

- 34. (Currently Amended) The method of claim 3129 or 32, wherein the indicator cell contains a recombinant expression vector encoding the human c-Maf protein.
- 35. (Currently Amended) The method of claim 3129 or 32, wherein the reporter gene is operatively linked to regulatory sequences of a Th2-associated cytokine gene.
- 36. (Previously Presented) The method of claim 34, wherein the human c-Maf-coding sequences are operatively linked to regulatory sequences that allow for constitutive expression of human c-Maf in the indicator cell.

37. (Currently Amended) The method of claim 34, wherein the human c-Maf coding sequences are operatively linked to regulatory sequences of the endogenous human c-Maf gene, wherein the regulatory sequences of the endogenous human c-Maf gene comprise the untranslated sequences of the Nhel/XbaI fragment of pHu-c-Maf (ATCC Accession No. 98671).

- 38. (Currently Amended) The method of claim 3129, wherein the reporter gene is a Th2-associated cytokine.
- 39. (Currently Amended) The method of claim <u>32 or 38</u>, wherein the Th2-associated cytokine is interleukin-4.
- 40. (Currently Amended) The method of claim 3129, wherein the reporter gene comprises nucleotides -157 tp to +58 relative to the +1start site of transcription of the proximal interleukin-4 promoter gene.
- 41. (Currently Amended) The method of claim 3129, wherein the reporter gene comprises about 3 kb of upstream regulatory sequences of the interleukin-4 gene.
- 42. (Currently Amended) The method of claim 3129, wherein the reporter gene is selected from the group consisting of genes that encode: chloramphenicol acetyltransferase, beta-galactosidase, alkaline phosphatase or and luciferase.

43. (Currently Amended) The method of claim 3429, wherein the indicator cell is derived from a cell line which does not normally express human c-Maf.

- 44. (Currently Amended) The method of claim 3129, wherein the indicator cell is derived from a B cell.
- 45. (Currently Amended) The method of claim 44, wherein the indicator cell is derived from the a M12 B lymphoma cell line.
- 46. (Currently Amended) The method of claim 3129, wherein the indicator cell is derived from a Th1 cell clone.
- 47. (Currently Amended) The method of claim 46, wherein the indicator cell is derived from an AE7 cells.
- 48. (Currently Amended) The method of claim 3129, wherein the indicator cell is a nonlymphoid cell.
- 49. (Previously Presented) The method of claim 48, wherein the indicator cell is a HEPG2 hepatoma cell.

50. (Previously Presented) The method of claim 48, wherein the indicator cell is a yeast cell.

- 51. (Currently Amended) The method of claim 32, wherein the effect of the test compound on an immune response is determined by determining the effect of the compound on expression production of a Th2-associated cytokine gene protein.
- 52. (Previously Presented) The method of claim 51, wherein the Th2-associated cytokine gene is an interleukin-4 gene.
- 53. (Currently Amended) The method of claim 32, wherein the effect of the test compound on an immune response the expression of a Th2-associated cytokine gene is determined by determining the effect of the compound on the development of T helper type 1 (Th1) cells.
- 54. (Currently Amended) The method of claim 32, wherein the effect of the test compound on an immune response the expression of a Th2-associated cytokine gene is determined by determining the effect of the compound on the development of T helper type 2 (Th2) cells.

55.-56. (Cancel)

57. (Currently Amended) A method for identifying a compound that modulates the activity of a human c-Maf protein comprising,

providing an indicator cell comprising a recombinant expression vector encoding

a human c-Maf protein comprising the Nhel/Xbal fragment of pHu-c-Maf (ATCC

Accession No. 98671), human c-Maf and a cytokine gene responsive to the human c-Maf protein,

contacting the indicator cell with the a test compound, and

determining the effect of the test compound on human c-Maf activity by

evaluating the level of cytokine production in the indicator cell in the presence and

absence of the test compound, wherein a modulation of the level of cytokine production

identifies the test compound as a modulator of the activity of a human c-Maf protein.

- 58. (Previously Presented) The method of claim 57, wherein the level of cytokine production is determined by detecting cytokine mRNA in the indicator cell.
- 59. (Previously Presented) The method of claim 57, wherein the level of cytokine production is determined by detecting cytokine secretion into the culture supernatant.
- 60. (Previously Presented) The method of claim 57, wherein the cytokine is interleukin-4.